

- I. Claim(s) 16 and 17 drawn to recombinant polypeptide comprising SEQ ID NO:2 or encoded by SEQ ID NO:1, classified in class 530, subclass 350.
- II. Claim(s) 16-17 drawn to recombinant polypeptide comprising SEQ ID NO:3 or encoded by SEQ ID NO:4, classified in class 530, subclass 350.
- III. Claim(s) 18-19, drawn to antibody which binds to recombinant polypeptide comprising SEQ ID NO:2 or encoded by SEQ ID NO:1, classified in class 530, subclass 387.9, for example.
- IV. Claim(s) 18-19, drawn to antibody which binds to recombinant polypeptide comprising SEQ ID NO:4 or encoded by SEQ ID NO:3, classified in class 530, subclass 387.9, for example.

In support of the present restriction requirement, the Examiner has alleged that the subject matter defined by the claims of the present invention represents four separate and distinct inventions. The Examiner alleges that "the products of Inventions I-IV are distinct because they have distinct functional, chemical and physical properties capable of separate use and manufacture".

As indicated, and in order to be fully responsive to the Examiner's requirements for restriction, Applicants provisionally elect to prosecute with traverse, the subject matter of Group IV, Claims 18-19, drawn to antibodies which bind to a recombinant polypeptide comprising SEQ ID NO:4 which is encoded by SEQ ID NO:3.

Pursuant to 37 C.F.R. §§1.111 and 1.143, Applicants hereby traverse the Examiner's requirement for restriction regarding Groups III and IV and request reconsideration thereof in view of the following remarks.

An Examiner's authority to require restriction is defined and limited by statute:

If two or more independent and distinct inventions are claimed in one application, the Commissioner may require the application to be restricted to one of the inventions.

35 U.S.C. §121, first sentence (emphasis added).

The implementing regulations of the Patent and Trademark Office include the mandate that restriction is appropriate only in cases presenting inventions which are both independent and distinct, 37 C.F.R. §§1.141-142. Without a showing of independence and distinctness, a restriction requirement is unauthorized.

In the present application, the claims of Groups III and IV which the Examiner has grouped separately are not "independent and distinct" so as to justify the restriction requirement. The antibodies of Groups III and IV are cross-reactive.

Accordingly, Groups III and IV are very clearly interrelated and interdependent, not "independent and distinct".

In addition, the courts have recognized that it is in the public interest to permit applicants to claim several aspects of their invention together in one application, as the Applicants have done herein. The CCPA has observed:

We believe the constitutional purpose of the patent system is promoted by encouraging applicants to claim, and therefore to describe in the manner required by 35 U.S.C. §112 all aspects as to what they regard as their invention, regardless of the number of statutory classes involved.

In re Kuehl, 456 F.2d 658, 666, 117 U.S.P.Q. 250, 256 (CCPA 1973). This interest is consistent

an invention customarily is sufficient to support claims in the same application to other aspects of the invention.

Applicants respectfully suggest that in view of the continued increase of official fees and the potential limitation of an applicants financial resources, a practice which arbitrarily imposes three-way restriction requirements may become prohibitive and thereby contravene the constitutional purpose to promote and encourage the progress of science and the useful arts. Moreover, under the regulatory changes as a consequence of the implementation of the General Agreement on Trade and Tariffs (GATT), Applicants are required either to conduct simultaneous prosecution, as here requiring excessive filing costs, or otherwise compromise the term of their patent assets.

It is vital to all applicants that restriction requirements issue only with the proper statutory authorization, because patents issuing on divisional applications which are filed to prosecute claims that the Examiner held to be independent and distinct can be vulnerable to legal challenges alleging double patenting. The third sentence of 35 U.S.C. §121, which states that a patent issuing on a parent application "shall not be used as a reference" against a divisional application or a patent issued thereon, does not provide comfort to applicants against such allegations. The Court of Appeals for the Federal Circuit has declined to hold that §121 protects a patentee from an allegation of same-invention double patenting, Studiengesellschaft Kohle mbH v. Northern Petrochemical Co., 784 F.2d 351, 355, 288 U.S.P.Q. 837, 840 (Fed. Cir. 1986); and in Gerber Garment Technology Inc. v. Lectra Systems Inc., 916 F.2d 683, 16 U.S.P.Q. 2d 1436 (Fed. Cir. 1990) that court held that §121 does not insulate a patentee from an allegation of "obviousness-patent that had issued from a divisional application filed following a restriction requirement



claims assigned to one group, on patent references found in the subclass(es) with which the Examiner associated another group of claims.

Furthermore, the classification system is a poor basis for requiring restriction between related aspects of an invention because classifications and definitions change over time. Thus, a classification that might have seemed to support restriction at a given time could change, thereby casting a shadow over the propriety of the restriction requirement later on during the term of the patents issuing from parent and divisional applications. Indeed, classifications seem largely to change in response to considerations of administrative convenience, and often in response to nothing more than growth in the number of patents in a given class or subclass. These considerations have nothing to do with whether the subject matter of patents assigned to different classifications is "independent and distinct" as those terms are used in 35 U.S.C. §121, which fact proves that basing restriction requirements on the classification system is improper.

In view of the foregoing comments, it is respectfully urged that the Examiner reconsider and withdraw the requirement for restriction regarding Groups III and IV and provide an action on the merits with respect to all the claims.

Respectfully submitted,



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